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සොම්ස පෝෂණ සහ දේශීය චෛදස අමාතසාංශය சுகாதார, போசணை மற்றும் சுதேச வைத்திய அமைச்சு Ministry of Health, Nutrition & Indigenous Medicine

General circulo 1-35 /2017

Provincial Directors of Health Services,

Regional Directors of Health Services,

Heads/ Directors of Institutions,

Directors of National Hospital/Teaching Hospitals/Provincial & District General Hospitals, Base Hospitals,

All Medical Superintendents of other Hospitals,

Heads of Decentralized units,

Provincial CCP,

All Regional Epidemiologists/ Medical Officers (Maternal and Child Health),

All Medical Officers of Health,

Measles, Rubella, Congenital Rubella Syndrome (CRS) elimination initiative - Sri Lanka
Measles is a highly infectious viral disease responsible for a high degree of morbidity and
mortality among children including complications of pneumonia (1-6%), diarrhoea (8%), Otitits
Media (7-9%), subacute sclerosing panencephalitis (SSPE) (1 per 100,000 cases), Keratitis and
Corneal scaring are common with Vitamine A deficiency.

Fatal cases of measles are now rarely reported in Sri Lanka after successful implementation of the National Immunization Programme, including 2 doses of measles, mumps and rubella (MMR) vaccination at 9 months and 3 years of age.

The measles vaccine was first introduced into the National Immunization Programme in Sri Lanka in 1984. Since then, morbidity and mortality of measles were reduced remarkably but outbreaks have been experienced in 1999-2000 and 2013-2015. Considering the requirement to enhance the population level immunity, 2nd dose of measles containing vaccine has been introduced with the measles, rubella (MR) vaccine in 2001. In 2011, MMR vaccine was introduced in 2 doses at the age of 1 year and at the age of 3 years, replacing measles (9 months) and MR (3 years) vaccines. But, considering the morbidity patterns and sero survey evidence during the measles outbreak situation in 2013-2015, the Advisory Committee on Communicable Diseases (ACCD) has decided to re-schedule the MMR 1st dose at 9 months of age, continuing the 2nd dose at 3 years.

Rubella is a mild disease affecting children and adults. However rubella in pregnant women is important as the virus is transmitted to the foetus across the placental barrier, sometimes with significant teratogenic effects. Rubella vaccine was introduced into the National Immunization Programme in 1996, targeting all reproductive age females of 11-44 years, with the objective of preventing congenital rubella syndrome (CRS). This was carried out as a school based programme by giving rubella vaccine to all children aged 11-15 years, and vaccinating the rest at the community clinics. Number of measles and CRS cases have markedly reduced and surveillance of measles, rubella and CRS was strengthened in 2005-2010 under the plan of 'intensification of the surveillance and Laboratory confirmation was made available for all suspected cases of Measles, Rubella and CRS from there to date.

Acceleration of measles, rubella, CRS Elimination Plan 2017-2020

In par with the Regional Measles, Rubella and CRS elimination strategic plans, Sri Lanka has set the goal of elimination of Measles, Rubella, CRS by 2020.

<u>Vision</u>: Sri Lanka is free from measles, rubella and CRS

Goal: To achieve and sustain measles, rubella and CRS free status in Sri Lanka

Objectives: To achieve and maintain zero endogenous transmission of measles, rubella and CRS in Sri Lanka and identify and contain possible imported outbreaks

Elimination targets:

- Zero endogenous measles cases by 2020
- Zero endogenous rubella cases by 2020
- Zero CRS case/ 100,000 live births by 2018

Components of elimination strategies:

- Achieve and maintain high levels of population immunity by providing two doses of measles and rubella containing vaccines with high vaccination coverage
- Strengthened disease surveillance including laboratory confirmation of all suspected cases of measles, rubella, CRS cases: case based investigation
- Strengthen country preparedness for outbreak prevention and response : contain
- Adequate patient care management to prevent the transmission and mortality
- Perform research to generate evidence for cost effective implementation strategies for

Measles and Rubella vaccination

 All eligible children who have completed the age of 9 months and the age of 3 years are to be vaccinated with MMR vaccine according to the current National Immunization schedule in Sri Lanka

- Required to achieve and maintain above 95% coverage in each of the two doses of MMR vaccine at the national, district and Medical Officer of Health (MOH) and Public Health Midwife (PHM) area levels
- If any child is found unvaccinated / missed for measles or rubella at any age, vaccinate with two doses of MMR with minimum of 6-8 weeks interval
- Ensure all women in the reproductive age are protected with at least one rubella containing vaccine (RCV)
- Ensure that that all women are protected/vaccinated for rubella at the time the Public Health Midwife (PHM) includes them in the Eligible Couple Register or at the earliest contact
- If any pregnant woman is found unvaccinated or with doubtful vaccination against rubella (and if the family has not been completed) she should be vaccinated with RVC after delivery, to prevent a future CRS case
- Once MMR (10 dose) vial is planned to open in the scheduled immunization clinic session and if the number of children planned for the days is less than the number in the opened vials for the day, plan and take necessary measures to vaccinate adults (up to 45 year) who are without proper history of measles and rubella vaccination using the remaining MMR doses for the day without discarding (after screen for contraindications and AEFI)

Surveillance Case definitions

Surveillance case definition of measles and rubella

Any person with "Fever and Maculopapular (i.e. non vescicular) rash" should be notified as either suspected measles or rubella case based on the clinical judgment of the treating clinicians / health care personnel

Surveillance case definition of CRS

Any infant with: maternal history of Rubella infection and / or with signs and symptoms from following categories

- cataract, congenital glaucoma, pigmentary retinopathy, congenital heart disease (PDA/peripheral pulmonary artery stenosis/VSD), Loss of hearing
- Purpura, splenomegaly, microcephaly, mental retardation, meningo-encephalitis, radiolucent bone disease, jaundice (within 24 hours of delivery)
- Laboratory data consistent with Congenital Rubella Infection (Rubella IgM positive or Rubella virus isolated)

Measles, Rubella, CRS case reporting

All suspected "measles and rubella" patients with "fever and maculopapular rash" should be notified by all medical officers who are treating the patient at first contact of the patient.

All other health care staff including field health staff, who meet with a patient of "fever and maculopapular rash" are required to inform to the immediate contact health authority for proper notification.

All hospitals where specialist paediatricians and / or physicians are available, are sentinel site hospitals for active surveillance for Measles/Rubella/CRS and weekly zero reporting.

- All suspected Measles and Rubella patients should be notified to the Epidemiology Unit through the updated 'Suspected Measles / Rubella Patient Information Form' (EPID/151/2/2015, Blue Form) [Annexure 1] filled by the Clinician/Medical Officer who is treating the patient at first patient contact.
- The routine notification should to be sent to the Medical Officer of Health (MOH) of the patient's residential area (Notification of Communicable Diseases: Health-544, Annexure 2) for all suspected cases of Measles, Rubella, CRS
- All <u>suspected CRS cases</u> need to be reported to the Epidemiology Unit immediately by phone/fax/e-mail and <u>special investigation form (EPID/DS/CRS/2013)</u> [Annexure 3] is required to be properly completed by the clinician/medical officer who is treating the patient at the health institution and to be sent to the Epidemiology Unit.
- All infection control nursing officers (ICNO) at the sentinel site hospitals are expected to
 maintain Measles/Rubella and CRS registers (Format: Annexure 4 and 5). The infection
 control nurses are also expected to visit medical, paediatric, obstetric, cardiology,
 ophthalmology and ENT wards regularly for detection of cases (all Measles, Rubella,
 CRS), actively look for cases and notify promptly to the Epidemiology Unit.
- All suspected cases of Measles, Rubella/CRS presented to sentinel site hospitals should be included in the Weekly reporting form for AFP, Measles, Rubella cases from hospital (sentinel sites) EPID/37/5/R2004 (Annexure 6) [or in the web based system which will be trained during the year], and should be completed for the week ending date of Friday and should be sent to the Chief Epidemiologist, Epidemiology Unit, Colombo with copy to the Regional Epidemiologist. This form should be sent even if no cases have been detected ("Nil" reporting) for the week. A total of 52 reports should be received from each site per year and the timeliness of the return needs to be maintained at 7 days to be received at the Epidemiology Unit. The performance rate of completeness and the timeliness of the return will be measured to maintain the surveillance performance.
- The patients identified in other health institutions including General Practitioners and private health care institutions, are required to be promptly notified to the relevant MOH (Notification of Communicable Disease, [Health 544] form or any other means of notification) and the laboratory confirmation should be carried out as instructed.
- The Medical Officer of Health of the Patients residence (in an institutional outbreak, the MOH of the institution belonged) has to proceed with the routine surveillance procedure, contact tracing and outbreak prevention for all notified or community detected Measles, Rubella, CRS cases and complete the special field investigation form for all clinically confirmed measles or rubella cases (irrespective of the laboratory confirmation or the availability of results.

Clinically confirmed measles case	Clinically confirmed rubella case		
Fever and maculopapular rash patient with at least one of the following: Cough Coryza (i.e. runny nose) Conjunctivitis (i.e. red eyes)	Fever with maculopapular rash and arthralgia, arthritis, lymphadenopathy (usually suboccipital/ postauricular/ cervical) or conjunctivitis		

- All clinically confirmed cases of Measles, Rubella and suspected CRS need to be completed with updated special investigation forms by the MOH ([EPID/DS/MEASLES/2007], [EPID/DS/RUBELLA.2007], [EPID/DS/CRS/2013]) (Annexure 7, 8 & 3) and duly completed forms should be sent to the Epidemiology unit as early as possible, maximum with 2 weeks delay from the date of the notification.
- If the notified/clinically confirmed measles/rubella case has not been tested for laboratory confirmation due to any reason by the health institution, the MOH should perform the laboratory testing at the time of special field investigation as per instructions in the Epidemiology Unit letter No: EPID/151/2011 dated 20/09/2012.
- All Measles/Rubella cases detected at the community level by any of the public health staff, need to be adequately investigated, in accordance with the routine surveillance and special investigation procedure, with laboratory testing procedure to complete case based investigation by the MOH.

Laboratory investigations for suspected Measles/Rubella and CRS cases

Two types of samples should be collected from all suspected measles and rubella cases

Sample for Virus isolation	Sample for detection of IgM (recent infection)
Nasal and throat swabs (in virus transport medium) preferably in the first 5 days of the onset of rash	2-3 ml blood sample preferably from 3 rd to 28 th day of the onset of rash

- A blood sample of 2-3ml for Measles /Rubella IgM should be collected from each suspected case of Measles, or Rubella from the 3rd day to 28th day of the onset of signs and symptoms, into a sterile, dry, screw capped container without any anti coagulant.
- A blood sample for Rubella IgM or for TORCH screen (as for Toxoplasma, Rubella, Cytomegalovirus, Herpes simplex virus in screening for congenital abnormalities) should be taken from all suspected infants of CRS and from newborns in instances where the mother has declared a history of suspected/confirmed Rubella infection in any gestational age of pregnancy.
- If any pregnant woman who does not give a history of Rubella vaccination is identified in any Obstetric Unit, she is required to be tested for Rubella IgM (before or after delivery) to identify possible recent Rubella infection during gestational period. In case a positive result is obtained, the baby is required to be investigated on delivery and followed up for possible CRS.

- Once the blood sample has been collected, it should be labeled and left at room temperature for about 30 minutes for clot formation. The sample should be sent as early as possible to the Measles and Rubella, National Reference Laboratory, Medical Research Institute (MRI), Colombo with a properly completed specimen request form (Annexure 9: "Specimen Request Form: Measles and Rubella, National Reference Laboratory, Medical Research Institute (MRI), Colombo). The sample should be transported in a cold box with ice cubes / ice packs to maintain cold temperature.
- If a facility to centrifuge is available, properly labelled separated serum should be sent to the MRI for Measles or Rubella IgM detection.
- The serum / clotted blood sample should be received at the laboratory within 48 hours of collection and if there is any delay of transport more than 6 hours the sample should be refrigerated until dispatch to prevent destruction of antibodies.
- Naso-pharyngeal aspirate, throat swab or gingival swab is collected within the first 5 days of the onset of symptoms for measles / rubella virus detection. Samples should be collected in to the container with virus transport medium (VTM) and labelled. Samples should be stored immediately at the refrigerator and transport in ice to maintain cold temperature with the completed specimen request form (annexure 9). Specimen collection containers (VTM + swabs) are provided. Contact infection control nursing officer (ICNO) of the hospital or Regional Epidemiologist of the district.

Measles, rubella outbreak response:

- Routine surveillance for outbreak detection and prevention after notification (initial Public health Inspector visit, field level investigation with Health H-411 / H-411a, and MOH case based special form investigation including laboratory confirmation should be completed within 14 days of the onset of the rash)
- Even a single laboratory confirmed measles, or rubella case is detected, it should be considered as an outbreak and following measures should be taken
 - Immediate notification to the Epidemiology Unit/ National focal point for Measles Rubella Elimination Programme(Epidemiology Unit)/RDHS/ Provincial CCP/ Regional Epidemiologist
 - If any unvaccinated/ unprotected child (2 doses of MMR or adult (up to 45 years) in the household, take measures to provide MMR vaccination at earliest possible, preferably within 14-21 days of the onset of the index laboratory confirmed case
 - Screen 30-50 households or households of 1 km radius around the index household, to identify any unvaccinated children below 15 years: take measures to vaccinate if any
 - Exclusion of the continuation of the outbreak:
 - Follow up contacts for 2 incubation period cycles (minimum of 28 days)
 - identify all "fever and maculopapular rash" cases from the area and send samples for laboratory testing (include into the surveillance system)
 - Inform to Epidemiology Unit the action taken

Additional information

- Unprotected travellers to measles or rubella (with unknown history or unvaccinated for measles and not contracted measles or rubella disease),
 - travelling to an endemic country for any of these diseases, are advised to vaccinate/receive at least one MMR dose, with a minimum of 1 month before the travel date, from the nearest MOH office
 - any unprotected traveller, returning from an endemic country, develops fever and rash within 14 to 21 days of the return should be considered as a possible imported case of measles or rubella and should be adequately investigated, to prevent community transmission
- Measles, Rubella vaccination and surveillance activities in disaster situations should be paid special attention, and should continue with routine immunization. Contact Measles, Rubella, CRS elimination programme at the Epidemiology Unit, Consultant CCP, Regional Epidemiologist, or the area MOH to assess the situation, and advise and actions for special vaccination campaigns and prevention of possible outbreaks
- Measles / Rubella outbreak prevention and response, specimen collection guidelines (including field level)[as per Epidemiology Unit letter No: EPID/151/2011 dated 20/09/2012], standard operation procedure (SOP) for specimen collection and transport, Accelerated measles, rubella, CRS elimination plan 2017-2020 are available in the website: http://www.epid.gov.lk, under disease information, Measles, Rubella, CRS elimination programme
- Additional information contact: Measles, Rubella, CRS Elimination Programme, Epidemiology Unit, No: 231, De Saram Place, Colombo 10, chepid@sltnet.lk, Tel:0112695112, fax: 0112696583

Please bring the contents of this circular to the notice of all relevant staff at your institution/district/province and arrange to implement the programme accordingly.

Dr. J.M.W. Jayasundara Bandara Director General of Health Services Ministry of Health Dr. J. M. W. Jayasundara Bandara
Director General of Health Services (Acting)
Ministry of Health, Nutrition & Indigenous Medicine,
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Colombo 10.

Copy:

- Secretary Health
- DDG/PHS 1
- DDG/PHS 11
- DDG/MS I and II
- DDG/Laboratory Services
- Chief Epidemiologist
- Director/ Private Healthcare Institutions
- Director/ MCH
- Director/FHB
- Director/NIHS
- Directors / Millitary hospitals

EPIDEMIOLOGY UNIT – MINISTRY OF HEALTH Measles / Rubella Elimination Initiative Suspected Measles / Rubella Patient Information

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List of Notifiable Diseases

ෑ" කාණ	ව ේචය	பிரிவு A		Group-A	
0	කොළරාව	9	வாந்திபேதி	9	Cholera
	මතාමාරිය		பிளேக் (கொள்ளை நோய்)	0	Plague
	කාභ උණ	۰	மஞ்சள் காய்ச்சல்	0	Yellow Fever
ත" කාද	න්ඩය ්	பிரிவு B		Group-B	
0	උගු බාලක පක්ෂගාත රෝගය පැපොල රෝගය		இளம்பிள்ளை வாதம் / சடுதியான தளர்ச்சி வாதம் கொப்பளிப்பான்	0	Acute Poliomyclitis / Acute Flacci Paralysis
0	ඩෙංගු උණ / ඩෙංගු රක්තපාත උණ	0		0	Chicken pox
0	ගලපටලය		டெங்கு காய்ச்சல் / டெங்கு குருதிப்பெருக்கு காய்ச்சல்		Dengue Fever / Dengue
-	රක්ත අතීසාරය		தொண்டைக் கரப்பன்		Haemorhagic Fever
0	නිදිකර්පථපුදාහය		வயிற்றோட்டம்		Diptheria
	අාත්තික උණ		முளைக்காய்ச்சல் (என்கெபலைடிஸ்)	9	Dysentary Encephalitis
0	ආහාර විෂවීම	6	நெருப்புக்காய்ச்சல் (குடல் காய்ச்சல்)		Enteric Fever
9	මානව ජලභීතිකා රෝගය	9	உணவு நச்சுத் தன்மை		Food poisoning
	ලෙප්ටොස්පයිරෝසියාව		விசர்விலங்குக்கடி நோய்		Human Rabies
	මැලේරියාව		லெப்டோஸ்பைரோசிஸ்		Leptospirosis
٠	සරම්ප	9	மலேரியா		Malaria
	මස්තිෂ්ක පටල පුදාහය		சின்னமுத்து		Measles
0	කම්මුල්ගාය		மூளைக்காய்ச்சல் (மெனின்சைடிஸ்)		Meningitis
	රුබෙල්ලා (ජර්මන් සරම්ප)	۰	கூகைக்கட்டு		Mumps
9	ස-ජාතීය රුබෙල්ලා රෝගය		ருபெல்லா/ருபெல்லா நோயுடன் பிறப்பு	a	Rubella / Congenital Rubella
0	සරල කල්පවත්තා උණ	۰	7 நாட்களுக்கு மேல் தொடரும்		Syndrom
	පිටගැස්ම		சாதாரண காய்ச்சல்		Simple Continued Fever of over
6	තවජත්ම පිටගැස්ම		ஏற்ப <mark>ுவ</mark> லி		7days or more
0	ටයිපස් උණ		பிறந்த முதல் மாதத்தில் ஏற்புவலி	8	Tetanus
9	වෛරස් යාකෘති පුදාහය		தைபசுக் காய்ச்சல்	•	Neonatal Tetanus
0	කක්කල් කැස්ස		வைரசு ஈரல் அழற்சி		Typhus Fever
٥	ක්ෂය රෝගය		குக்கல்	0	Viral Hepatitis
1750			காச நோய்	•	Whooping Cough Tuberculosis
0	ලීෂ්මනයිසිස් -		லேயி'மேனியாசிஸ்	0	Leishmaniasis
9	ලාදුරු		தொழுநோய		Leprosy

කරුණාකර මෙතතින් නවන්න / இங்கே மடிக்கவும் / Please Fold Here

රා. සේ. පි. / அரச சேவை / O. S. S සෞඛා වෛදා නිලධාරි சுகாதார வைத்திய அதிகாரி

THE MEDICAL OFFICER OF HEALTH

කරුණාකර මෙතතින් අලවන්න / இங்கே ඉட்டவும் / Please Paste Here

EPID/DS/CRS/2013

SURVEILLANCE OF CONGENITAL RUBELLA SYNDROME (CRS) - CASE INVESTIGATION FORM EPIDEMIOLOGY UNIT, MINISTRY OF HEALTH

The Medical Officer/Hospital and REE/MOH should carry out the investigation personally. Necessary data should be obtained from the mother of the new baby/BHT/Physician/investigation reports/diagnosis cards. Early investigation and return is essential.

A CENEDAL		Serial No:	
A. GENERAL 1. Date of notification to MOH: 2. Date of notification to Epidemio 3. Name of the reporting Institution 4. Ward No: 5. BHT No: 6. Name of the hospital where the 7. Ward No:	logy Unit:	(dd/mm/yy)	
8. BHT No:			
9. Name of patient (BLOCK LETT 10. Name of the parent/guardian 11. Residential Address:	ERS)		
12. Date of Birth :	15. Ethnic group	16. Mothers occupa	ation
B. PRESENT ILLNESS /OUTC 19. Date of detection of signs and symptoms of CRS: d d mm y y 20. Where did the patient detect first 1. Government hospital 2. Private hospital 3. Medical Officer of Health 4. Private practitioner 5. Ayurvedic institution 6. Other (specify)	21. Outcome of the event 1. Still under treatment 2. Died 3. Transferred 4. Discharged 22. Date of discharge, trans (where relevant) d d m m y y	24. sfer or death	If transferred, name of hospital Was patient transferred from some other hospital Yes / No / If "yes", where was the patient transferred from ?
. CLINICAL DATA urveillance Case definition: hild <1 year of age with maternal hi	story of Rubella infection and/or fo	ollowing signs and sympto	oms.
List A 1. Cateract/s 2. Congenital glaucoma 3. Congenital hearth disease 4. Loss of hearing 5. Pigmentary Retinopathy	List B 1. Purpura 2. Splenomegaly 3. Microcephaly 4. Mental Retardation 5. Meningo-encephalitis 6. Radiolucent bone disease 7. Jaundice	Laboratory data consismith Congenital Rubell Infection (CRI) positive result of rubella IgM	For office use only Compatible with the case definition. 1. Yes 2. No

Investigation	Date of collection of specimen (dd/mm/yy)	Laboratory MRI/ other govt./ private/ not known	Results (mark MA if ter results are not available and PP if pending)
1. maternal IgG persisting >6/52 in infant		TIOT KHOWN	and PF ii pending)
2. rubella specific IgM			
3. virus isolation / PCR			
. MATERNAL HISTORY	F. MOTHER'S IMMUNIZATION		ACT HISTORY
Age of mother at time of delivery:	34. Was the mother immunized for	1 00. 1100 110	mother in contact with a know
years		3. not known or susp index p	ected case of rubella during tregnancy?
0. Did the mother have a rubella-like illness	35. If yes, date of vaccination:	1 1.5	/es
during the present pregnancy?			
1. yes	d d m m y y		
2. no	not known	3. r	not known
3. not known	36. Type of vaccine used:	40. If yes, pe	eriod of gestation in weeks:
 If yes, period of gestation at the time of illness 	1. Rubella 3. MR		7
in weeks	37. Place of vaccination	not	known
not known	1. MOH clinic		
2. Which of the following symptoms and	2. school		
signs were present?	3. government hospital		
1. fever	4. private dispensary/surge	ery .	
2. rash	5. private hospital		
3. lymphadenopathy	6. other (specify)		
4. conjunctivitis	7. not known		
5. arthritis/arthralgia		-	
6. others (specify)	38. If not immunized, reason:		
	2. unaware of the need for		
Was rubella serologically confirmed	3. non-availability of vaccin		
during pregnancy?	4. no faith in the vaccine	4 14 14 14	
1. yes	5. others (specify)	Time b	FOR OFFICE USE etween immunization and
2. no		develop	ment of maternal infection
3. not known	6. not known		yrs months
Remarks:			
nature:	Name:	The second of th	For office use only
nature:	Name:	Acres de la companya	For office use only
		Final c	classification
nature:	Name: Designation:	Final c	

Measles / Rubella Register Format

					ø			
							Serial N (Annual r	
							Dise	ase
							Patient	Name of the
							DOB	Δρρ
							Sex	
							OPD	Ward
							OPD	
							Date of ad	mission
							Date of ons	
							dose	MMR/MR/Mea sles/ Rubella
							dose	nation MR/Mea es/ pella
							Residential address	
							To MOH (H 544)	Dat notifi
						Y	To Epid Unit (EPID/151/1/2013)	Date of notification
							Blood for IgM	collec confii (da re
			2				Virus isolation swabs	Specimen collection for confirmation (date & result)

CRS Register Format

	8				Serial Number (Annual number)	
					Name of the Patient & Name of the mother	
					Age at detection of CRS & DOB	
					Sex	
					Ward/ Unit	9
					внт	
					Date of admission	
					Date of detection	
		i.			Mother's vaccination status Rubella (Y/N)	
					Residential address	
					To MOH (H 544) notification	Date of
					To Epid Unit (EPID/DS/CRS/2013)	of
	27				Blood for IgM (date & result) Virus isolation swabs	Specimen collection
					Virus isolation swabs mation	collection

FORM: EPID/37/5/R2004

WEEKLY REPORTING FORM FOR AFP*, MEASLES, RUBELLA /CRS CASES FROM HOSPITALS

(SENTINEL SITES)

Disease	Name of the patient	Age	Sex	Ward	B.H.T. No.	** D:O.A	Date of onset	Residential address
5		9						
					# # # # # # # # # # # # # # # # # # #			
teamainteam.organipadiatikasijaasija					·			
paediatric his returi copy to R	c. EYE, ENT of every Frida egional Epida ettnet.lk by Ho	ind neur y to Epi emiologi ead of th	ology w demiolo ist, Tel: ie institi	ards dur gist, Epic 2695112 ution/IC	ing the we demiologi , 2681548 N/PHI or	ek. Even i cal Unit, i, Fax: 26 any other	f no cases 231, de Sa 96583, E-n identified	FORM: EPID/37/5/R200
	WEEKLY	REPOF	RTING	FORM	HOSP	TTALS EL SITES	· K	RUBELLA/CRS CASES FROM
NSTITUTI	ION:	***********	, ,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	•	HOSP (SENTIN	TTALS EL SITES)	
NSTITUTI		***********	, ,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	•	HOSP (SENTIN	TTALS)	.200 Residential address
NSTITUTI	reporting: (S	Saturday	y to Frie	day) [HOSP (SENTIN	TTALS EL SITES	Date of	.200
NSTITUTI	reporting: (S	Saturday	y to Frie	day) [HOSP (SENTIN	TTALS EL SITES	Date of	.200
NSTITUTI	reporting: (S	Saturday	y to Frie	day) [HOSP (SENTIN	TTALS EL SITES	Date of	.200

EPID/DS/MEASLES/2016

SURVEILLANCE OF MEASLES - CASE INVESTIGATION FORM EPIDEMIOLOGY UNIT MINISTRY OF HEALTH

The MOH should	l do the investigation perso	onally. Necessary	y data should be obtained fro	m the hospital by reference to the BHT/Physician or
from the diagnos	sis card. Early investigation	and return is es	ssential.	
Week ending of Notification	d d m m y y	Date of Confirmation		ID Register No : Mea

		-		
Week ending d d m m y	Date of Confirmation	d d m m y y	ID Register No : Mea	/
A. PARTICULARS OF PATIEN	IT (Please (✓) app	propriate box where	applicable)	
Name of patient (BLOCK LETTE) Residential Address:	RS)			
	***************************************	email :		
3. Date of Birth :/ / 4. Age	LILI (dd/mm/yy)			
□□/□□ □ 1. Male	6. Ethnic group 7 ☐ 1. Sinhalese	. Occupation	8. RDHS area	9. MOH area
y y / m m 2. Female	2. Tamil			
y y / mm	3. Moor	***************************************		
	9. Unknown			
P DDECENT II I NECO (OUTO)				
B. PRESENT ILLNESS /OUTCO				
To. (a) Date of onset of fever	12. Was patient a	dmitted to hospital?	17. Outcome of	the case
	☐ 1. Yes ☐ 2. No	(If "Yes" question 13) (If "No" skip to question	☐ 1. C	
(b) Date of onset of rash	2.140	(II INO Skip to question		
(5) Bate of offset of fash	13. If yes, date of	admission:		omplication parge, transfer or death
			To. Date of discr	Tarister or death
d d m m y y	d d m m	уу	d d m	m yy
11. (a) Did the patient seek medical	14 Name of heavil		10 If transferred	
advice 1. Yes 2. No	14. Name of nospit	tal:		name of hospital
(b) If "Yes" where did the patient first seek medical advice?	15. Ward :			
1. Government hospital			20. Was patient t	ransferred from some
2. Private hospital	16. BHT No:		other hospita	
3. Private practitioner			1. Yes 2. No	
4. Ayurvedic institution				
(public/private)			21. If "Yes", when	e was the patient
5. Other (specify)			transferred from	

C. CLINICAL DATA			hiterary organizations in the	
Case definition: fever and maculop	apular rash with one	of cough, coryza (run	ny nose) or conjunctivitie	
22. Symptoms and signs Yes No				
	,	23. Compli	cations Yes N	0
1. fever]	1. none		
2. masculopapular rash		2. diarrhoea]
3. cough		pneumor		
5. conjunctivitis		4. otitis med]
6. other (Specify)		5. encephal 6. other (sp]
		o. other (sp	ecny)	
). LABORATORY FINDINGS				
4. Was blood taken for measles serolo	ogy (measles IgM) ?	1. Yes 2.	No 🗆	
a) If yes: 1. Hospital 🗌 2. P	rivate Practitioner	3. MOH 4. Othe	r 🔲	
0)				

Investigation (Serology) e.g. IgM / IgG	Date of collection of specimen (dd/mm/yy)	Date of sent to laboratory	Laboratory (MRI/govt./private)	Results (mark NA if test results are not available)	Date of results

San	mple type		Date of	Date of sent		Name of the		Date of result	ult		Result	S
			collection			laborato				+ve	-ve	Ge
☐ (i) swabs (th ☐ (ii) secretion ☐ (iii) urine ☐ (iv) other												typ
. MEASLES	VACCINA	TION STAT	TUS									
6. Was Measle	s Containing	g Vaccine giv	ven (MCV) [Mea		MMR]							
			1	Yes 🗌	2. N	0	3.1	Not known				
27. If "yes" (a) Number of doses		er of doses	1 🔲		2 🔲		>2	>2 🗌				
	(b) source	of information	on V	accination	card 🔲	☐ History: ☐		tory:				
etails of immun	ization											
ose		Date of im	munization		f vaccine	MD.	Batch	number		Place	1	
st dose		(SG/THTI/Y)		ivieasie	es, MR, MI	VIT				mmur	nization	×
nd dose									-			
ther												0
1011 40 11			Hospital / Priva	0.00								
1. Yes [(if yes, f	2. ill row 1 – 3	No with details)	nyone with feve 3. Not know	vn 🔲								
1. Yes [(if yes, f	2. ill row 1 – 3 ient's house	No Uwith details)		vn 🔲	loped a sin					nt of ı	measle	es in
1. Yes [(if yes, f	2. ill row 1 – 3 ient's house	No Uwith details)	3. Not know	vn 🔲	loped a sins)		s <u>follov</u>	ving the dev	elopme		measle	
1. Yes [(if yes, f	2. ill row 1 – 3 ient's house their immun	No Uwith details)	3. Not know close contacts s (fill Row 4 – 7	who devel	loped a sin s)	nilar illnes	s <u>follo</u> v	ving the dev	elopme		for MC	V Not
1. Yes [(if yes, f etails of the pat e patient, and a. contacts with	2. ill row 1 – 3 ient's house their immun	No Uwith details)	3. Not know close contacts s (fill Row 4 – 7	who devel	loped a sins)	nilar illnes	s <u>follov</u>	ving the dev	elopme	nated	for MC	V Not
1. Yes [(if yes, f stails of the pat e patient, and a. contacts with imilar disease or to onset of	2. ill row 1 – 3 ient's house their immun	No Uwith details)	3. Not know close contacts s (fill Row 4 – 7	who devel	loped a sins)	nilar illnes	s <u>follov</u>	ving the dev	elopme	nated	for MC	V Not
1. Yes [(if yes, f stails of the pat e patient, and a. contacts with imilar disease or to onset of ess in the	2. ill row 1 – 3 ient's house their immun	No Uwith details)	3. Not know close contacts s (fill Row 4 – 7	who devel	loped a sins)	nilar illnes	s <u>follov</u>	ving the dev	elopme	nated	for MC	V Not
1. Yes [(if yes, f) etails of the pate patient, and a. contacts with imilar disease or to onset of ess in the ient b. contacts who	2. ill row 1 – 3 ient's house their immun Name	No Uwith details)	3. Not know close contacts s (fill Row 4 – 7	who devel	loped a sins)	nilar illnes	s <u>follov</u>	ving the dev	elopme	nated	for MC	V Not
1. Yes [(if yes, f) etails of the pate patient, and a. contacts with imilar disease or to onset of ess in the ient b. contacts who reloped similar ess after	2. ill row 1 – 3 ient's house their immun Name	No Uwith details)	3. Not know close contacts s (fill Row 4 – 7	who devel	loped a sins)	nilar illnes	s <u>follov</u>	ving the dev	elopme	nated	for MC	V Not
1. Yes [(if yes, f etails of the pat e patient, and a. contacts with imilar disease or to onset of ess in the itent c. contacts who veloped similar ess after ntact of the	2. iill row 1 – 3 ient's house their immun Name 1 2 3 4	No Uwith details)	3. Not know close contacts s (fill Row 4 – 7	who devel	loped a sins)	nilar illnes	s <u>follov</u>	ving the dev	elopme	nated	for MC	V Not
1. Yes [(if yes, f etails of the pat e patient, and a. contacts with imilar disease or to onset of ess in the itent c. contacts who veloped similar ess after ntact of the	2. ill row 1 – 3 ient's house their immun Name 1 2 3 4 5	No Uwith details)	3. Not know close contacts s (fill Row 4 – 7	who devel	loped a sins)	nilar illnes	s <u>follov</u>	ving the dev	elopme	nated	for MC	V Not
1. Yes [(if yes, f) etails of the patie patient, and a. contacts with similar disease or to onset of less in the tient b. contacts who veloped similar ess after nact of the lex patient	2. iill row 1 – 3 ient's house their immun Name 1	No □ with details) hold or other ization status	3. Not know	who devel with details	loped a sir s) Date of rash	nilar illnes	s <u>follov</u>	ving the dev	elopme	nated	for MC	V Not
1. Yes [(if yes, f) etails of the pate patient, and a. contacts with similar disease or to onset of ess in the tient b. contacts who veloped similar ess after ntact of the ex patient	2. iill row 1 – 3 ient's house their immun Name 1	No □ with details) hold or other ization status	3. Not know close contacts s (fill Row 4 – 7	who devel with details	loped a sir s) Date of rash	nilar illnes	s <u>follov</u>	ving the dev	elopme	nated	for MC	V Not
1. Yes [(if yes, f) etails of the pate patient, and a. contacts with imilar disease or to onset of ess in the ient b. contacts who veloped similar ess after patient list the p	2. iill row 1 – 3 ient's house their immun Name 1	No with details) hold or other ization status	3. Not know	who devel with detail. Sex	Date of rash	nilar illness fonset of	Relat patie	ving the dev	Vaccin Yes	nated	for MC	V Not
1. Yes [(if yes, f) etails of the patie patient, and a. contacts with imilar disease or to onset of ess in the ient b. contacts who reloped similar ess after lact of the ex patient Is the patient I es: (i) Count	2. iill row 1 – 3 ient's house their immun Name 1	No with details) hold or other ization status	3. Not know close contacts s (fill Row 4 – 7 Age	who devel with detail: Sex Sex	Date of rash	onset of	Relat patie	ving the dev	Vaccin Yes	nated	for MC	V Not
1. Yes [(if yes, f) etails of the patie patient, and a. contacts with imilar disease or to onset of ess in the ient b. contacts who reloped similar ess after lact of the ex patient Is the patient I es: (i) Count	2. iill row 1 – 3 ient's house their immun Name 1	No with details) hold or other ization status	3. Not know close contacts s (fill Row 4 – 7 Age	who devel with detail. Sex	Date of rash	onset of	Relat patie	ving the dev	Vaccin Yes	nated	for MC	V Not
1. Yes [(if yes, f) etails of the pate patient, and a. contacts with similar disease or to onset of ess in the tient b. contacts who veloped similar ess after stact of the ex patient Is the patient I es: (i) Count (ii) Histor	2. iill row 1 – 3 ient's house their immun Name 1	No with details) hold or other ization status	3. Not know close contacts s (fill Row 4 – 7 Age	who devel with detail: Sex Sex	Date of rash	onset of	Relat patie	ving the dev	Vaccin Yes	nated	for MC	V Not
1. Yes [(if yes, f) etails of the pate patient, and a. contacts with similar disease or to onset of ess in the tient b. contacts who veloped similar ess after thact of the ex patient a. Is the patient I ess: (i) Count (ii) Histor	2. ill row 1 – 3 ient's house their immun Name 1 2 3 4 5 6 7 naving a histry of travel:	No hold or other ization status	3. Not know close contacts s (fill Row 4 – 7 Age	who devel with detail: Sex Sex	loped a sirs) Date of rash Ilness ons	et)	Relation	ving the dev	Vaccii Yes	nated No	for MC	VVNot
1. Yes [(if yes, f) etails of the pat e patient, and a. contacts with similar disease or to onset of less in the tient b. contacts who veloped similar ess after ntact of the lex patient . Is the patient if less: (i) Count	2. ill row 1 – 3 ient's house their immun Name 1 2 3 4 5 6 7 naving a histry of travel:	No hold or other ization status	3. Not know	who devel with detail: Sex Sex	loped a sirs) Date of rash Ilness ons	et)	Relation	ving the devisionship to nt	Vaccii Yes	nated No	for MC	VVNot
1. Yes [(if yes, f) etails of the pate patient, and a. contacts with imilar disease or to onset of ess in the ient b. contacts who veloped similar ess after plact of the ex patient Is the patient I les: (i) Count (ii) Histor	2. iill row 1 – 3 ient's house their immun Name 1	No with details) hold or other ization status tory of travel	3. Not know close contacts s (fill Row 4 – 7 Age abroad (3 weel	who devel with detail: Sex	loped a sins) Date of rash Ilness ons	et)	Relation	ving the devisionship to nt	Vaccii Yes	nated No	for MC	V
1. Yes [(if yes, f) etails of the patie patient, and a. contacts with similar disease or to onset of eass in the tient b. contacts who veloped similar ess after ntact of the lex patient Is the patient lex (ii) Count (iii) Histor marks	2. ill row 1 – 3 ient's house their immun Name 1 2 3 4 5 6 7 naving a histry of travel:	No with details) hold or other ization status tory of travel s contact in a	3. Not know	who devel with detail: Sex Sex	loped a sirs) Date of rash Ilness ons	et)	Relation	ving the devisionship to nt	Vaccii Yes No R OFFIC Labor	nated No	for MC	V Not know

EPID/DS/RUBELLA/2007

... 1. Yes 2. No

SURVEILLANCE OF RUBELLA - CASE INVESTIGATION FORM

EPIDEMIOLOGY UNIT, MINISTRY OF HEALTH

The MOH should do the investigation personally. Necessary data should be obtained from the hospital by reference to the BHT / Physician or from the diagnosis card. Early investigation and return are essential. Please write the Serial No given in the Infectious Disease Register (ID Register) in the MOH office of notification Serial no: A. PARTICULARS OF PATIENT (Please tick (✓) the appropriate box where applicable) Name of patient (BLOCK LETTERS) 2. Residential address: 3. Date of birth: (dd/mm/yyyy) 4. Age 5. Sex 6. Ethnic group 7. Occupation 8. DPDHS division (district) 9. MOH area 1. male 1. Sinhalese 2. female 2. Tamil y/m m 3. not known 3. Moor FOR OFFICE USE ONLY 4. others ☐5. not known B. PRESENT ILLNESS/OUTCOME 10. Date of onset of symptoms: 12. Was patient admitted to hospital? 17. Date of discharge/transfer or death: → to Q. 13 1. yes skip 11. Where did the patient first seek 18. If transferred, name of hospital to Q. 21 3. not known medical advice? 13. If yes, date of admission: 1. government hospital 19. Was patient transferred from some other hospital? 2. private hospital d m 1. yes ____ 2. no 3. private practitioner 14. Name of hospital: 20. If "yes", where was the patient transferred from? 4. Ayurvedic institution (public/private) 21. Outcome of the case 15. Ward: 5. other (specify) 1. cured 3. transferred 16. BHT no: 2. died 4. not known C. CLINICAL DATA Case definition: An illness with generalized macular papular rash, fever and arthralgia/arthritis, lymphadenopathy or conjunctivitis 22. Symptoms and signs 23. Complications 1. fever 1. encephalitis 2. rash 2. other (specify): 3. lymphadenopathy For office use only 4. conjunctivitis Compatible with the 5. arthritis/arthralgia case definition: 6. other (specify):

D. LA	BORATORY	FINDINGS				
24. Was	s blood taken fo	or measles serology? 1. y	/es	not known		
5. If ye	es,					
Investigation		Date of collection of specimen (dd/mm/yy)	Labora (MRI/ other go not kno	vt./ private/	Results (mark NA if test results are n available and PP if pending)	
1. lgG	1 st specimen					
. IgG	2 nd specimen					
. IgM						
. Virus	s isolation		-			
6. Was		110000000000000000000000000000000000000	onset of the present illn	ess?		
	Dose	Date of immunization* (dd/mm/yy)	Type of vaccine**	Batch number	Place of immunization***	
	1 st dose					
	2 nd dose					
	Other	s not known but the particular dos				
Was t EXP Was t If yes,	yes ☐ 2. no OSURE DUR the patient preg , period of gest	raccine 5. not known	emales of reproduc	☐ 6. other (spe and rash) in the ma tive age only)	onth prior to the onset of rash?	
pregn S case Rema	ant mothers we investigation	vho had an acute attack sho n form No EPID/DS/CRS/200	ould be followed up. If 7 must be filled.	the baby is foun	d to have acquired CRS, a separate	
nature:			Name:		For office use only Final classification Laboratory confirmed	
demio		niology Unit, 231, De Saram		2696583	Epidemiologically confirmed	

Specimen Request Form: Measles & Rubella National Reference Laboratory Medical Research Institute, Colombo

	Name of the Patient	•		
•	Age :		Sex -:-	
•	Hospital	:		
•	Ward	:		•
•	BHT No:	:		
•	Address of patient's residence	:		
			***************************************	***************************************
•	District of patient's residence	:		
•	Date of onset of fever	:	DD MM YYYY	
•	Date of onset of rash	:	GD MM YYYY	
•	Specimen collection	:	Blood	Throat swab
			Date of collection	Date of collection
Ě			DD MM YYYY	DU MM YYYY
	*:		(3 ml blood preferably serum separated & transport in cold box)	(throat swabs in virus transport media in Ice)
•	Date of dispatch to MRI	:	DD MM YYYY	
*Ple EPI	ase notify the case to <u>Medical</u> D/151/1/2013 – Blue Form)	of	icer of health (H 544)	& to Epidemiology Unit

	e a		Signature of M	edical Officer
	LA	BC	RATORY USE	
	Date received of sample	:	Serology	Virus Isolation
			DD MW AAAA	DO MM YYYY
	Sample accepted	:	Yes / No	Yes / No
	(Received within 72 hrs after collection	on,	sent in cold chain, satisfactor	ry sample)